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Serial No. 09/469,717

ETH 1623

1-46. (Cancelled)

- (Previously Amended) The anastomosis device of claim 60 wherein the tubular 47. member is pre-shaped and has at least a first bend along a length of the member.
- 48. (Previously Amended) The anastomosis device of claim 60wherein a portion of the tubular member extends at an angle of between 30° and 90° relative to a longitudinal centerline.
- 49. (Previously Amended) The anastomosis device of claim 60wherein said tubular member is formed from a biocompatible material.
- 50. (Original) The anastomosis device of claim 49 wherein said biocompatible material is biocrodable.
- (Previously Amended) The anastomosis device of claim 49 wherein said 51. biocompatible material comprises a polymeric material.
- 52. The anastomosis device of claim 51 wherein said polymeric material is (Original) selected from a group consisting of a polymer, a homopolymer, and a copolymer.
- 53. (Original) The anastomosis device of claim 52, wherein the polymeric material is a polycaprolactone.
- 54. (Previously Amended) The anastomosis device of claim 60wherein an end portion of the graft lumen is everted over an end margin of the tubular member.
- 55. (Previously Amended) The anastomosis device of claim 54 wherein the tubular member has an adhesive surface and the end portion of the graft lumen is adhered to the tubular member.

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Serial No. 09/469,717

ETH 1623

- 56. (Original) The anastomosis device of claim 49 wherein the tubular member includes a chromophore.
- 57. (Original) The anastomosis device of claim 56 wherein said chromophore is a dye.
- 58. (Original) The anastomosis device of claim 60wherein said tubular member is impregnated with one or more agents selected from the group consisting of anti-platelet, anti-thrombus, and anti-inflammatory compounds.
- 59. (Previously Amended) The anastomosis device of claim 60wherein the tubular member is impregnated with one or-more anti-proliferative compounds.
- 60. (Previously Amended) A fastener for sealingly joining a graft lumen to a target vessel in an anastomosis, the target vessel having an opening formed in a side wall thereof, the fastener comprising a tubular member formed of a deformable material and sized and dimensioned for receiving an end portion of said graft lumen, said tubular member being transformable upon application of energy to the tubular member between a fluent state in which the tubular member is radially expandable to permit radial expansion of the graft lumen vessel, and a non-fluent state in which the tubular member retains the end portion of the graft lumen in its expanded state in sealing engagement with the target vessel.

61-62. (Cancelled)

63. (Previously Amended) The fastener of claim 60 wherein the tubular member comprises a material selected from a group consisting of polyethylene-glycol (PEG) based hydrogels, acrylates, and acrylated urethanes.

64-67. (Cancelled)